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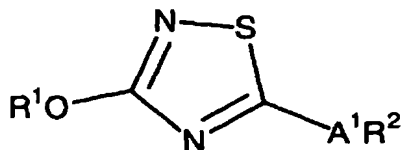
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(54) Title: 1,2,4-THIADIAZOLE COMPOUNDS AND ARTHROPOD CONTROLLING COMPOSITION CONTAINING THE
SAME



(1)

1,2,4-thiadiazole compound has an excellent arthropod controlling activity, and can effectively control an arthropod pests such
as insect pests, acarine pests and the like.

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DESCRIPTION

1,2,4-THIADIAZOLE COMPOUNDS AND ARTHROPOD CONTROLLING COMPOSITION CONTAINING THE SAME

TECHNICAL FIELD

5 The present invention relates to 1,2,4-thiadiazole compounds and arthropod controlling composition containing the same.

BACKGROUND ART

10 While various pesticide compositions have been used for the purpose of controlling arthropod pests such as insect pests, acarine pests and the like, sometimes the effect of those pesticide compositions is not always enough, and therefore the development of novel arthropod controlling compositions having enough effect is desired.

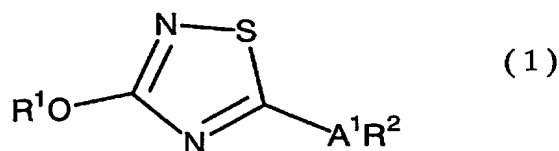
15 It is an objective of the present invention to provide novel compounds having a arthropod controlling effect, arthropod controlling compositions containing the same compounds, and methods for controlling arthropods applying effective dose of the same compounds to pests or their habitat.

DISCLOSURE OF INVENTION

20 The present inventors have intensively studied to find compounds having excellent arthropod controlling activity, and as a result, found out that the 1,2,4-thiadiazole compounds of formula (1) as depicted below have an excellent controlling activity for arthropod pests such as insect pests and acarine pests, thereby completing the present invention.

25 Namely, the present invention relates to a 1,2,4-thiadiazole

compound represented by the formula (1) (hereinafter referred to as the present invention compound(s)):



wherein, R¹ represents C3-C7 alkynyl that may be substituted with
5 halogen atom; R² represents C3-C8 cycloalkyl, C5-C8 cycloalkenyl or
C7-C11 bicycloalkyl, wherein the C3-C8 cycloalkyl, the C5-C8
cycloalkenyl and the C7-C11 bicycloalkyl may be substituted with a
substituent(s) selected from a group of C1-C4 alkyl, halogen atom
and trifluoromethyl; A¹ represents a single bond, C1-C2 alkylene or
10 C2-C3 alkylidene;

an arthropod controlling composition containing the present invention
compound as active ingredient; and a method for controlling arthropod pests
comprising applying an effective dose of the present invention compound to
arthropod pests or habitat of arthropod pests.

15

MODE FOR CARRYING OUT THE INVENTION

In the present invention, the following groups are exemplified as each
substituent described above.

The C3-C7 alkynyl that may be substituted with halogen atom
20 represented by R¹ includes, for example, 2-propynyl, 2-butynyl, 4-fluoro-2-
butynyl, 3-butynyl, 1-methyl-2-butynyl, 2-pentynyl, 3-pentynyl, 4,4-
dimethyl-2-pentynyl, 3-chloro-2-propynyl, 3-bromo-2-propynyl, 3-iodo-2-
propynyl and 1-methyl-2-propynyl.

In the C3-C8 cycloalkyl, the C5-C8 cycloalkenyl and C7-C11 bicycloalkyl represented by R², C1 to C4 alkyl as substituents includes, for example, methyl, ethyl, propyl, isopropyl and 1,1-dimethylethyl.

The C3-C8 cycloalkyl that may be substituted with a substituent(s) selected from a group of C1-C4 alkyl, halogen atom and trifluoromethyl represented by R² includes, for example, cyclobutyl, cyclopentyl, 2-methylcyclopentyl, 3-methylcyclopentyl, 2-ethylcyclopentyl, 3-ethylcyclopentyl, 2,5-dimethylcyclopentyl, 3,4-dimethylcyclopentyl, cyclohexyl, 2-methylcyclohexyl, 3-methylcyclohexyl, 4-methylcyclohexyl, 2-ethylcyclohexyl, 3-ethylcyclohexyl, 4-ethylcyclohexyl, 2-propylcyclohexyl, 3-propylcyclohexyl, 4-propylcyclohexyl, 3-isopropylcyclohexyl, 3,3-dimethylcyclohexyl, 3,5-dimethylcyclohexyl, 3,5-diethylcyclohexyl, 4,4-dimethylcyclohexyl, 3,3,5,5-tetramethylcyclohexyl, 3-fluorocyclohexyl, 3-chlorocyclohexyl, 3-bromocyclohexyl, 3-trifluoromethylcyclohexyl, 3,5-difluorocyclohexyl, 3,5-dichlorocyclohexyl, 3,5-dibromocyclohexyl, 3,5-bis(trifluoromethyl)cyclohexyl, 1-chlorocyclohexyl, 1-bromocyclohexyl, 1-chloro-3-methylcyclohexyl, 1-bromo-3-methylcyclohexyl, 1-chloro-3,5-dimethylcyclohexyl, 1-bromo-3,5-dimethylcyclohexyl, cycloheptyl, 2-methylcycloheptyl, 3-methylcycloheptyl, and cyclooctyl.

The C5-C8 cycloalkenyl that may be substituted with a substituent(s) selected from a group of C1-C4 alkyl, halogen atom and trifluoromethyl represented by R² includes, for example, 1-cyclopentenyl, 2-methyl-1-cyclopentenyl, 3-methyl-1-cyclopentenyl, 4-methyl-1-cyclopentenyl, 5-methyl-1-cyclopentenyl, 2-cyclopentenyl, 2-methyl-2-cyclopentenyl, 3-methyl-2-cyclopentenyl, 4-methyl-2-cyclopentenyl, 5-methyl-2-cyclopentenyl, 3-

cyclopentenyl, 2-methyl-3-cyclopentenyl, 3-methyl-3-cyclopentenyl, 4-methyl-3-cyclopentenyl, 5-methyl-3-cyclopentenyl, 1-cyclohexenyl, 2-methyl-1-cyclohexenyl, 3-methyl-1-cyclohexenyl, 4-methyl-1-cyclohexenyl, 5-methyl-1-cyclohexenyl, 6-methyl-1-cyclohexenyl, 2-cyclohexenyl, 2-methyl-2-cyclohexenyl, 3-methyl-2-cyclohexenyl, 4-methyl-2-cyclohexenyl, 5-methyl-2-cyclohexenyl, 6-methyl-2-cyclohexenyl, 3-cyclohexenyl, 2-methyl-3-cyclohexenyl, 3-methyl-3-cyclohexenyl, 4-methyl-3-cyclohexenyl, 5-methyl-3-cyclohexenyl, 6-methyl-3-cyclohexenyl, 1-cycloheptenyl, 2-cycloheptenyl, 3-cycloheptenyl, 1-cyclooctenyl, 2-cyclooctenyl, 3-cyclooctenyl and 4-cyclooctenyl.

The C7-C11 bicycloalkyl that may be substituted with a substituent(s) selected from a group of C1-C4 alkyl, halogen atom and trifluoromethyl represented by R^2 includes, for example, bicyclo[2.2.1]-2-heptyl, 7,7-dimethylbicyclo[2.2.1]-2-heptyl, bicyclo[2.2.1]-7-heptyl, bicyclo[2.2.2]-2-octyl, bicyclo[3.2.1]-2-octyl, *cis*-bicyclo[4.4.0]-2-decanyl and *trans*-bicyclo[4.4.0]-2-decanyl.

The C1-C2 alkylene represented by A^1 is methylene ($-\text{CH}_2-$) or ethylene ($-\text{CH}_2\text{CH}_2-$); the C2-C3 alkylidene represented by A^1 is ethylidene ($-\text{CH}(\text{CH}_3)-$), propylidene ($-\text{CH}(\text{CH}_2\text{CH}_3)-$) or isopropylidene ($-\text{C}(\text{CH}_3)_2-$).

As the aspects of the present invention compound, for example, the following compounds are exemplified.

a 1,2,4-thiadiazole compound wherein R^1 is 2-butyryl in the formula (1);

a 1,2,4-thiadiazole compound wherein R^1 is 2-pentyryl in the formula

(1);

a 1,2,4-thiadiazole compound wherein R² is cyclopentyl that may be substituted with C1-C4 alkyl in the formula (1);

5 a 1,2,4-thiadiazole compound wherein R² is cyclohexyl that may be substituted with C1-C4 alkyl in the formula (1);

a 1,2,4-thiadiazole compound wherein A¹ is single bond in the formula (1);

a 1,2,4-thiadiazole compound wherein A¹ is methylene in the formula (1);

10 a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl and R² is cyclopentyl that may be substituted with C1-C4 alkyl in the formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl and R² is cyclohexyl that may be substituted with C1-C4 alkyl in the formula (1);

15 a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl and R² is cyclopentyl that may be substituted with C1-C4 alkyl in the formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl and R² is cyclohexyl that may be substituted with C1-C4 alkyl in the formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl and A¹ is single bond that may be substituted with C1-C4 alkyl in the formula (1);

20 a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl and A¹ is methylene that may be substituted with C1-C4 alkyl in the formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl and A¹ is single bond that may be substituted with C1-C4 alkyl in the formula (1);

25 a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl and A¹ is methylene that may be substituted with C1-C4 alkyl in the formula (1);

a 1,2,4-thiadiazole compound wherein R² is cyclopentyl that may be substituted with C1-C4 alkyl and A¹ is single bond in the formula (1);

a 1,2,4-thiadiazole compound wherein R² is cyclopentyl that may be substituted with C1-C4 alkyl and A¹ is methylene in the formula (1);

5 a 1,2,4-thiadiazole compound wherein R² is cyclohexyl that may be substituted with C1-C4 alkyl and A¹ is single bond in the formula (1);

a 1,2,4-thiadiazole compound wherein R² is cyclohexyl that may be substituted with C1-C4 alkyl and A¹ is methylene in the formula (1);

10 a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl, R² is cyclopentyl that may be substituted with C1-C4 alkyl and A¹ is single bond in the formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl, R² is cyclohexyl that may be substituted with C1-C4 alkyl and A¹ is single bond in the formula (1);

15 a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl, R² is cyclopentyl that may be substituted with C1-C4 alkyl and A¹ is single bond in the formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl, R² is cyclohexyl that may be substituted with C1-C4 alkyl and A¹ is single bond in the formula (1);

20 a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl, R² is cyclopentyl that may be substituted with C1-C4 alkyl and A¹ is methylene in the formula (1);

25 a 1,2,4-thiadiazole compound wherein R¹ is 2-butynyl, R² is cyclohexyl that may be substituted with C1-C4 alkyl and A¹ is methylene in the

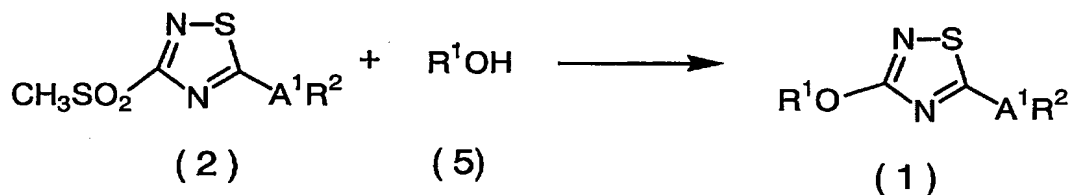
formula (1);

a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl, R² is cyclopentyl that may be substituted with C1-C4 alkyl and A¹ is methylene in the formula (1);

5 a 1,2,4-thiadiazole compound wherein R¹ is 2-pentynyl, R² is cyclohexyl that may be substituted with C1-C4 alkyl and A¹ is methylene in the formula (1).

10 The following will describe a production process for the present invention compounds.

The present invention compounds can be produced, for example, by reacting a sulfone compound (2) and an alcohol compound (5);



wherein R¹, R² and A¹ are as defined above.

15 The reaction is generally carried out in the presence of base in a solvent.

The solvent to be used in the reaction includes, for example, ethers such as 1,2-dimethoxyethane, 1,4-dioxane, tetrahydrofuran and the like, acid amides such as N,N-dimethylformamide and the like, and mixtures
20 thereof.

The base to be used in the reaction includes, for example, inorganic base such as sodium hydride, potassium carbonate and the like.

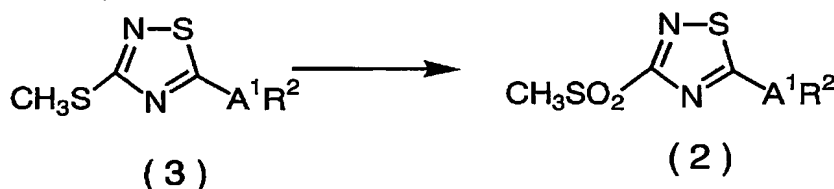
The amount of the base to be used in the reaction is usually 1 to 2

moles, and the amount of the alcohol compound (5) is usually 1 to 1.5 moles; relative to 1 mole of the sulfone compound (2).

The reaction temperature is usually in the range of -20°C to 80°C, and the reaction time is usually in the range of 1 to 24 hours.

5 After completion of the reaction, the present invention compound can be isolated by subjecting the reaction mixture to ordinary post-treatment such as adding the reaction mixture into water, extracting with an organic solvent, concentrating the organic phase obtained and the like. The isolated present invention compound can be purified by a technique such as
10 chromatography and the like, if necessary.

The sulfone compound (2) can be produced by reacting a sulfide compound (3) and an oxidizing agent;



15 wherein R² and A¹ are as defined above.

The reaction is generally carried out in a solvent.

The solvent to be used in the reaction includes, for example, halogenated hydrocarbons such as chloroform, dichloromethane and the like.

The oxidizing agent to be used in the reaction includes, for example,
20 peroxyacetic acid, 3-chloroperoxybenzoic acid and the like.

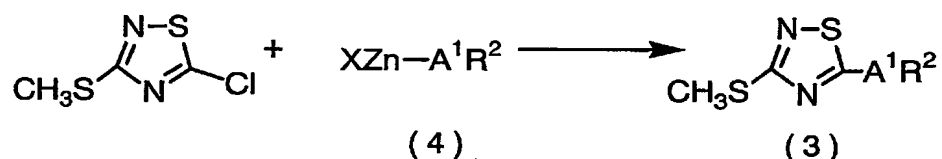
The amount of the oxidizing agent to be used in the reaction is usually 2 to 2.5 moles relative to 1 mole of the sulfide compound (3).

The reaction temperature is usually in the range of -20°C to 60°C,

and the reaction time is usually in the range of 1 to 24 hours.

After completion of the reaction, the sulfone compound (2) can be isolated by subjecting the reaction mixture to ordinary post-treatment such as adding the reaction mixture into aqueous solution of sodium
 5 hydrogensulfite, extracting with an organic solvent, concentrating the organic phase obtained and the like. The isolated sulfone compound (2) can be purified by a technique such as chromatography and the like, if necessary.

The sulfide compound (3) can be produced by reacting 5-chloro-3-methylthio-1,2,4-thiadiazole and an organozinc compound (4) in the
 10 presence of transition metal catalyst;



wherein R² and A¹ are as defined above, and X represents bromine atom or iodine atom.

15 The reaction is generally carried out under atmosphere of inactive gas such as nitrogen and argon, in a solvent.

The solvent to be used in the reaction includes, for example, ethers such as 1,2-dimethoxyethane, 1,4-dioxane, tetrahydrofuran and the like, aromatic hydrocarbon such as benzene, toluene and the like.

20 The transition metal catalyst to be used in the reaction includes, for example, palladium catalyst such as palladium(II) acetate, tetrakis(triphenylphosphine) palladium(0), {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane

complex and bis(triphenylphosphine) palladium(II) chloride.

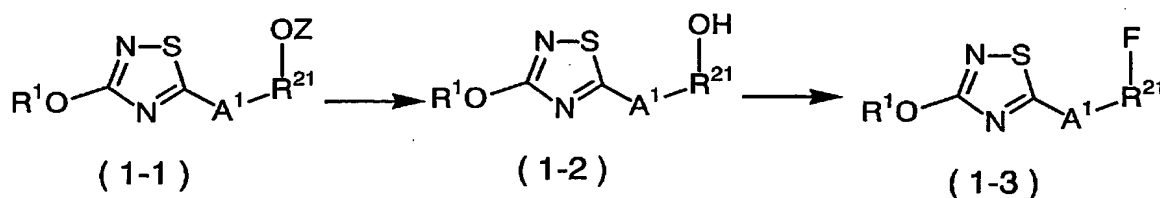
The amount of the transition metal catalyst is usually 0.001 to 0.1 mole, and the amount of the organozinc compound (4) is usually 0.9 to 1.5 moles; relative to 1 mole of 5-chloro-3-methylthio-1,2,4-thiadiazole.

5 The reaction temperature is usually in the range of 0 to 150°C, and the reaction time is usually in the range of 1 to 24 hours.

After completion of the reaction, the sulfide compound (3) can be isolated by subjecting the reaction mixture to ordinary post-treatment such as concentrating the reaction mixture as such, subjecting the residue to the chromatography and the like.

The organozinc compound (4) can be produced by reacting a halogenated compound represented with R^2A^1-X and metal zinc.

A part of the present invention compounds can be produced by a following method.

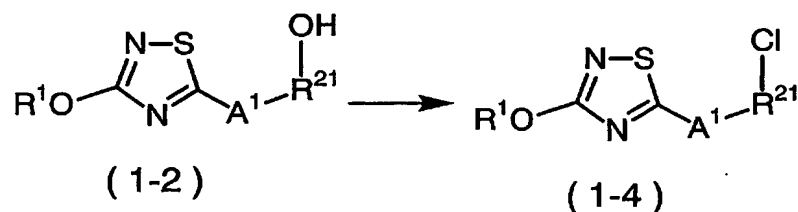


wherein R^1 and A^1 are as defined above, Z represents a protecting group of hydroxy such as methoxymethyl, and R^{21} represents C3-C8 cycloalkandiyl that may be substituted C1-C4 alkyl, C5-C8 cycloalkendiyl that may be substituted C1-C4 alkyl or C7-C11 bicycloalkandiyl that may be substituted C1-C4 alkyl.

The compound (1-2) can be produced by deprotecting of the protecting

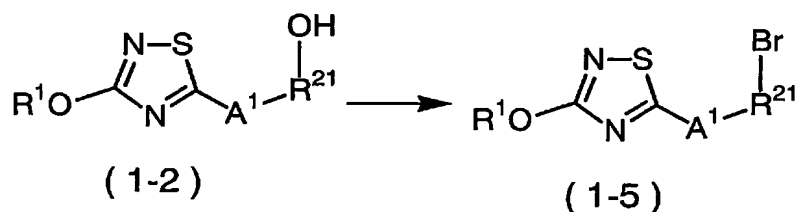
group of hydroxy of the compound (1-1). In a case of the protecting group represented by Z is methoxymethyl, for example, it can be produced by reacting with water in the presence of acid. The acid to be used in the reaction includes, for example, hydrochloric acid, sulfuric acid and paratoluenesulfonic acid.

The compound (1-3) can be produced by reacting the compound (1-2) and a fluorination reagent. The fluorination reagent to be used includes, for example, diethylaminosulfur trifluoride.



wherein R^1 , A^1 and R^{21} are defined above.

The compound (1-4) can be produced by reacting the compound (1-2) in the presence of triphenylphosphine and carbon tetrachloride.



wherein R^1 , A^1 , Z and R^{21} are defined above.

The compound (1-5) can be produced by reacting the compound (1-2) and a bromination reagent. The bromination reagent to be used includes, for example, triphenylphosphine bromide.

The examples of the present invention compound, which can be produced according to the above-described production processes, are listed below.

- 5 A 1,2,4-thiadiazole compound (from I.01.01 to I.01.89) wherein R¹ is 2-propynyl, A¹ is single bond and R² is a substituent selected from the group described below in the formula (1);
- a 1,2,4-thiadiazole compound (from I.02.01 to I.02.89) wherein R¹ is 2-butynyl, A¹ is single bond and R² is a substituent selected from the group
10 described below in the formula (1);
- a 1,2,4-thiadiazole compound (from I.03.01 to I.03.89) wherein R¹ is 1-methyl-2-butynyl, A¹ is single bond and R² is a substituent selected from the group described below in the formula (1);
- a 1,2,4-thiadiazole compound (from I.04.01 to I.04.89) wherein R¹ is 2-pentynyl, A¹ is single bond and R² is a substituent selected from the group
15 described below in the formula (1);
- a 1,2,4-thiadiazole compound (from I.05.01 to I.05.89) wherein R¹ is 2-butynyl, A¹ is methylene and R² is a substituent selected from the group described below in the formula (1);
- 20 a 1,2,4-thiadiazole compound (from I.06.01 to I.06.89) wherein R¹ is 1-methyl-2-butynyl, A¹ is single bond and R² is a substituent selected from the group described below in the formula (1);
- a 1,2,4-thiadiazole compound (from I.07.01 to I.07.89) wherein R¹ is 2-pentynyl, A¹ is single bond and R² is a substituent selected from the group
25 described below in the formula (1).

- cyclobutyl(01), cyclopentyl(02), 2-methylcyclopentyl(03), 3-methylcyclopentyl(04), 2-ethylcyclopentyl(05), 3-ethylcyclopentyl(06), 2,5-dimethylcyclopentyl(07), 3,4-dimethylcyclopentyl(08), cyclohexyl(09), 2-methylcyclohexyl(10), 3-methylcyclohexyl(11), 4-methylcyclohexyl(12), 2-ethylcyclohexyl(13), 3-ethylcyclohexyl(14), 4-ethylcyclohexyl(15), 2-propylcyclohexyl(16), 3-propylcyclohexyl(17), 4-propylcyclohexyl(18), 3-isopropylcyclohexyl(19), 3,3-dimethylcyclohexyl(20), 3,5-dimethylcyclohexyl(21), 3,5-diethylcyclohexyl(22), 4,4-dimethylcyclohexyl(23), 3,3,5,5-tetramethylcyclohexyl(24), 3-fluorocyclohexyl(25), 3-chlorocyclohexyl(26), 3-bromocyclohexyl(27), 3-trifluoromethylcyclohexyl(28), 3,5-difluorocyclohexyl(29), 3,5-dichlorocyclohexyl(30), 3,5-dibromocyclohexyl(31), 3,5-bis(trifluoromethyl)cyclohexyl(32), 1-chlorocyclohexyl(33), 1-bromocyclohexyl(34), 1-chloro-3-methylcyclohexyl(35), 1-bromo-3-methylcyclohexyl(36), 1-chloro-3,5-dimethylcyclohexyl(37), 1-bromo-3,5-dimethylcyclohexyl(38), cycloheptyl(39), 2-methylcycloheptyl(40), 3-methylcycloheptyl(41), cyclooctyl(42), 1-cyclopentenyl(43), 2-methyl-1-cyclopentenyl(44), 3-methyl-1-cyclopentenyl(45), 4-methyl-1-cyclopentenyl(46), 5-methyl-1-cyclopentenyl(47), 2-cyclopentenyl(48), 2-methyl-2-cyclopentenyl(49), 3-methyl-2-cyclopentenyl(50), 4-methyl-2-cyclopentenyl(51), 5-methyl-2-cyclopentenyl(52), 3-cyclopentenyl(53), 2-methyl-3-cyclopentenyl(54), 3-methyl-3-

cyclopentenyl(55), 4-methyl-3-cyclopentenyl(56), 5-methyl-3-cyclopentenyl(57), 1-cyclohexenyl(58), 2-methyl-1-cyclohexenyl(59), 3-methyl-1-cyclohexenyl(60), 4-methyl-1-cyclohexenyl(61), 5-methyl-1-cyclohexenyl(62), 6-methyl-1-cyclohexenyl(63), 2-cyclohexenyl(64), 2-methyl-2-cyclohexenyl(65), 3-methyl-2-cyclohexenyl(66), 4-methyl-2-cyclohexenyl(67), 5-methyl-2-cyclohexenyl(68), 6-methyl-2-cyclohexenyl(69), 3-cyclohexenyl(70), 2-methyl-3-cyclohexenyl(71), 3-methyl-3-cyclohexenyl(72), 4-methyl-3-cyclohexenyl(73), 5-methyl-3-cyclohexenyl(74), 6-methyl-3-cyclohexenyl(75), 1-cycloheptenyl(76), 2-cycloheptenyl(77), 3-cycloheptenyl(78), 1-cyclooctenyl(79), 2-cyclooctenyl(80), 3-cyclooctenyl(81), 4-cyclooctenyl(82), bicyclo[2.2.1]-2-heptyl(83), 7,7-dimethylbicyclo[2.2.1]-2-heptyl(84), bicyclo[2.2.1]-7-heptyl(85), bicyclo[2.2.2]-2-octyl(86), bicyclo[3.2.1]-2-octyl(87), *cis*-bicyclo[4.4.0]-2-decanyl(88) and *trans*-bicyclo[4.4.0]-2-decanyl(89).

The arthropod pests against which the present invention compound has control activity may include, for example, insect pests and acarine pests. Specific examples are listed below:

20

Hemiptera: Delphacidae such as *Laodelphax striatellus*, *Nilaparvata lugens*, and *Sogatella furcifera*; Deltocephalidae such as *Nephotettix cincticeps* and *Empoasca onukii*; Aphididae such as *Aphis gossypii* and *Myzus persicae*; Pentatomidae; Aleyrodidae such as *Trialeurodes vaporariorum*, *Bemisia tabaci*, and *Bemisia argentifolii*; Coccidae; Tingidae;

25

Psyllidae;

Lepidoptera: Pyralidae such as *Chilo suppressalis*, *Cnaphalocrocis medinalis*, *Ostrinia nubilalis*, and *Parapediasia teterrella*; Noctuidae such as *Spodoptera litura*, *Spodoptera exigua*, *Pseudaletia separata*, *Mamestra brassicae*, *Agrotis ipsilon*, *Thoricoplusia* spp., *Heliothis* spp., *Helicoverpa* spp., and *Earias* spp.; Pieridae such as *Pieris rapae crucivora*; Tortricidae such as *Adoxophyes orana fasciata*, *Grapholita molesta*, and *Cydia pomonella*; Carposinidae such as *Carposina niponensis*; Lyonetiidae such as *Lyonetia clerkella*; Gracillariidae such as *Phyllonorycter ringoniella*;
 10 Phyllocnistidae such as *Phyllocnistis citrella*; Yponomeutidae such as *Plutela xylostella*; Gelechiidae such as *Pectinophora gossypiella*; Arctiidae; Tineidae;

Diptera: Calicidae such as *Culex pipiens pallens*, *Culex tritaeniorhynchus*, and *Culex quinquefasciatus*; Aedes spp. such as *Aedes aegypti* and *Aedes albopictus*; Anopheles spp. such as *Anopheles sinensis*;
 15 Chironomidae; Muscidae such as *Musca domestica* and *Muscina stabulans*; Calliphoridae; Sarcophagidae; Fanniidae; Anthomyiidae such as *Delia platura* and *Delia antiqua*; Tephritidae; Drosophilidae; Psychodidae; Tabanidae; Simuliidae; Stomoxyidae; Agromyzidae;

20 Coleoptera: Diabrotica spp. such as *Diabrotica virgifera virgifera* and *Diabrotica undecimpunctata howardi*; Scarabaeidae such as *Anomala cuprea* and *Anomala rufocuprea*; Curculionidae such as *Sitophilus zeamais*, *Lissorhoptrus oryzophilus*, and *Callosobruchus chienensis*; Tenebrionidae such as *Tenebrio molitor* and *Tribolium castaneum*; Chrysomelidae such as
 25 *Oulema oryzae*, *Aulacophora femoralis*, *Phyllotreta striolata*, and

Leptinotarsa decemlineata; Anobiidae; Epilachna spp. such as *Epilachna vigintioctopunctata*; Lyctidae; Bostrychidae; Cerambycidae; *Paederus fuscipes*;

Thysanoptera: Thripidae spp. including Thrips spp. such as *Thrips palmi*, Frankliniella spp. such as *Frankliniella occidentalis*, and Sciltothrips spp. such as *Sciltothrips dorsalis*; Phlaeothripidae spp.;

Hymenoptera: Tenthredinidae; Formicidae; Vespidae;

Dictyoptera: Periplaneta spp.; Blatta spp.;

Orthoptera: Acrididae; Gryllotalpidae;

10 Aphaniptera: Pulex irritans;

Anoplura: Pediculus humanus;

Isoptera: Termitidae;

Acarina: Tetranychidae.

15 The arthropod controlling composition of the present invention contains the present invention compound and an inert carrier. Generally, it is a preparation obtained by mixing the present invention compound and a carrier such as a solid carrier, a liquid carrier and a gaseous carrier, and if necessary, adding a surfactant and other adjuvant for formulation. The
20 formulation includes, for example, an emulsion, an oil solution, a shampoo formulation, a flowable formulation, a powder, a wettable powder, a granule, a paste formulation, a microcapsule, a foam, an aerosol, a carbon dioxide gas formulation, a tablet, a resin formulation and the like. These formulations can be converted to use into a poison bait, a pesticide coil, an electric
25 pesticide mat, a smoking agent, a fumigant or sheet.

In the pesticide composition of the present invention, the present invention compound is usually contained in an amount of 0.1% to 95% by weight.

The solid carrier for formulation includes, for example, a fine power
5 and a granule of clays (e.g., kaolin clay, diatomite, bentonite, Fubasami clay, acid clay, etc.), synthetic hydrated silicon oxide, talc, ceramic, other inorganic minerals (e.g., sericite, quartz, sulfur, activated carbon, calcium carbonate, hydrated silica) or chemical fertilizers (e.g., ammonium sulfate, ammonium phosphate, ammonium nitrate, ammonium chloride, urea).

10 The liquid carrier for formulation includes, for example, aromatic or aliphatic hydrocarbons (e.g., xylene, toluene, alkyl naphthalene, phenylxylylethane, kerosine, light oil, hexane, cyclohexane), halogenated hydrocarbons (e.g., chlorobenzene, dichloromethane, dichloroethane, trichloroethane), alcohols (e.g., methanol, ethanol, isopropyl alcohol, butanol,
15 hexanol, ethylene glycol), ethers (e.g., diethyl ether, ethylene glycol dimethyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, propylene glycol monomethyl ether, tetrahydrofuran, dioxane), esters (e.g., ethyl acetate, butyl acetate), ketones (e.g., acetone, methyl ethyl ketone, methyl isobutyl ketone, cyclohexanone), nitriles (e.g.,
20 acetonitrile, isobutyronitrile), sulfoxides (e.g., dimethylsulfoxide), acid amides (e.g., N,N-dimethylformamide, N,N-dimethylacetamide), vegetable oils (e.g., soy bean oil, cotton seed oil), vegetable essential oils (e.g., orange oil, hyssop oil, lemon oil) and water.

The gaseous carrier for formulation includes, for example, butane gas,
25 chlorofluorocarbons, liquefied petroleum gas (LPG), dimethyl ether, carbon

dioxide and the like.

The surfactant for formulation includes, for example, alkyl sulfate salts, alkylsulfonic acid salts, alkylarylsulfonic acid salts, alkyl aryl ethers and their polyoxyethylene derivatives, polyethylene glycol ethers, polyhydric
5 alcohol esters, and sugar alcohol derivatives.

The other adjuvant for formulation includes, for example, binders, dispersants and stabilizers, and specifically for example, casein, gelatin, polysaccharides (e.g., starch, gum arabic, cellulose derivatives, alginic acid), lignin derivatives, bentonite, sugars, synthetic water-soluble polymers (e.g.,
10 polyvinyl alcohol, polyvinylpyrrolidone, polyacrylic acid), PAP (isopropyl acid phosphate), BHT (2,6-di-t-butyl-4-methylphenol), BHA (a mixture of 2-t-butyl-4-methoxyphenol and 3-t-butyl-4-methoxyphenol), vegetable oils, mineral oils, fatty acids, and fatty acid esters.

A base for the poison bait includes, for example, grain powders,
15 vegetable oils, sugars, and crystalline cellulose, and further, if necessary, antioxidants such as dibutylhydroxytoluene and nordihydroguaiaretic acid, preservatives such as dehydroacetic acid, agents for preventing children and pets from erroneously eating such as hot pepper powder, and pest-attractive flavors such as cheese flavor, onion flavor and peanut oil may be added to
20 the base.

Arthropod pests can be controlled by applying an effective dose of the present invention compound to pests directly and/or habitats of pests (e.g., nest, plant, soil). Usually the preparation of the pesticide composition of the
25 present invention is used as the present invention compound.

When the pesticide composition of the present invention is used for a control of pests in agriculture and forestry, the application amount is usually 0.1 to 1,000 g as an active ingredient per 1000 m². The emulsions, wettable powders, flowables and microcapsule formulations are usually applied after dilution with water to have an active ingredient concentration of 0.01 to 10000 ppm, while oil solution, powders and granules are usually applied as such. These preparations may be sprayed directly to the plant to be protected from pests. The pests living in a soil can be controlled by treating the soil with these preparations, and the preparations can also be applied to treat seedbeds prior to the planting plants or to treat planting holes or plant bottoms in the planting. Furthermore, the sheet preparation of the pesticide composition of the present invention can be applied by a method such as winding around plants, stretching in the vicinity of plants and laying on the soil surface at the plant bottom.

When the pesticide composition of the present invention is used for a control of epidemic, the application amount is usually 0.001 to 10 mg as an active ingredient per 1 m³ in case of application for open space, and 0.001 to 100 mg as an active ingredient per 1 m² in case of application for plane surface. The emulsions, wettable powders and flowables are usually applied after dilution with water to have an active ingredient concentration of 0.01 to 100,000 ppm, while oil solutions, aerosols, smoking agents and poison baits are usually applied as such, and pesticide coils and electric pesticide mats are applied with emitting active ingredients by heating depending on their formulation form.

The pesticide composition of the present invention can also be used in admixture or combination with other insecticides, nematocides, acaricides, fungicides, herbicides, plant growth regulators, synergists, fertilizers, soil conditioners, animal feeds, and the like.

5 The active ingredients of such other insecticide and acaricide include, for example, pyrethroid compounds such as allethrin, tetramethrin, prallethrin, phenothrin, resmethrin, cyphenothrin, permethrin, cypermethrin, alpha-cypermethrin, zeta-cypermethrin, deltamethrin, tralomethrin, cyfluthrin, beta-cyfluthrin, cyhalothrin, lambda-cyhalothrin, 10 flumethrin, imiprothrin, etofenprox, fenvalerate, esfenvalerate, fenpropathrin, silafluofen, bifenthrin, transfluthrin, flucythrinate, tau-fluvalinate, acrinathrin and tefluthrin; organophosphorus compounds such as dichlorvos, fenitrothion, cyanophos, profenofos, sulprofos, phenthoate, isoxathion, tetrachlorvinphos, fenthion, chlorpyrifos, diazinon, acephate, 15 terbufos, phorate, chlorethoxyfos, fosthiazate, ethoprophos, cadusafos and methidathion; carbamate compounds such as propoxur, carbaryl, metoxadiazon, fenobucarb, methomyl, thiodicarb, alanycarb, benfuracarb, oxamyl, aldicarb and methiocarb; benzoylphenylurea compounds such as lufenuron, chlorfluazuron, hexaflumuron, diflubenzuron, triflumuron, 20 teflubenzuron, flufenoxuron, fluazuron, novaluron and triazuron; juvenile hormone-like substances such as pyriproxyfen, methoprene, hydroprene and fenoxycarb; neonicotinoid compounds such as acetamiprid, nitenpyram, thiacloprid, thiamethoxam and dinotefuran; N-phenyl-pyrazole compounds such as acetoprole and ethiprole; benzoylhydrazine compounds such as 25 tebufenozide, chromafenozide, methoxyfenozide and halofenozide;

diafenthiuron; pymetrozine; flonicamid; triazamate; buprofezin; spinosad;
emamectin benzoate; chlorfenapyr; indoxacarb MP; pyridalyl; cyromazine;
fenpyroximate; tebufenpyrad; tolfenpyrad; pyridaben; pyrimidifen;
fluacrypyrim; etoxazole; fenazaquin; acequinocyl; hexythiazox; clofentezine;
5 fenbutatin oxide; dicofol, propargite; abamectin; milbemectin; amitraz;
cartap; bensultap; thiocyclam; endosulfan; spiroidiclofen; spiromesifen; and
azadirachtin.

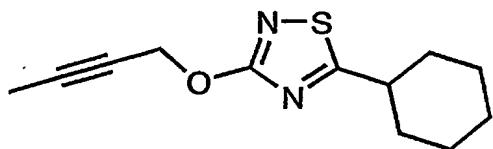
The present invention will be further illustrated by the following
10 production examples, formulation examples, and test examples; however,
the present invention is not limited to these examples. First, production
examples of the present invention compounds are exemplified.

In the following production examples, the reference production
examples, the data of ^1H -NMR were measured in a solvent of deuterium
15 chloroform with tetramethylsilane as the internal standard.

Production Example 1

200 mg of 3-methylsulfonyl-5-cyclohexyl-1,2,4-thiadiazole and 63 mg
of 2-butyne-1-ol were dissolved in 2 g of N,N-dimethylformamide, 39 mg of
sodium hydride (60% in oil) was added thereto under ice-cooling, and the
20 reaction mixture was stirred for 30 minutes under ice-cooling and for 1 hour
at room temperature. Then, the reaction mixture was added to saturated
brine, and extracted with t-butyl methyl ether. The organic layer was
concentrated under reduced pressure, and the residue obtained was
subjected to silica gel column chromatography to give 79 mg of 5-cyclohexyl-
25 3-(2-butyneoxy)-1,2,4-thiadiazole (hereinafter referred to as the present

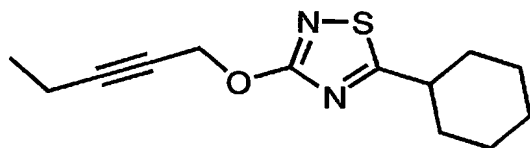
invention compound (1)).



$^1\text{H-NMR}$: 4.97 (s, 2H), 2.99 (m, 1H), 2.15 (d, 2H), 1.88 (m, 5H), 1.73 (d, 1H), 1.56-1.21 (m, 5H)

5 Production Example 2

By using 75 mg of 2-pentyn-1-ol instead of 2-butyn-1-ol according to Production Example 1 was obtained 98 mg of 5-cyclohexyl-3-(2-pentynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (2)).



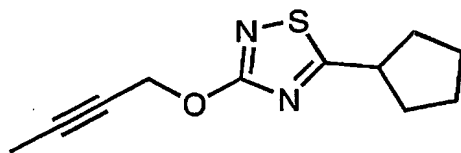
10

$^1\text{H-NMR}$: 5.00 (s, 2H), 2.99 (m, 1H), 2.24 (q, 2H), 2.13 (d, 2H), 1.85 (d, 2H), 2.74 (d, 1H), 1.56-1.24 (m, 5H), 1.14 (t, 3H)

Production Example 3

300 mg of 3-methylsulfonyl-5-cyclopentyl-1,2,4-thiadiazole and 99 mg
15 of 2-butyn-1-ol were dissolved in 3 g of N,N-dimethylformamide, 62 mg of sodium hydride (60% in oil) was added thereto under ice-cooling, and the reaction mixture was stirred for 30 minutes under ice-cooling. Then, the reaction mixture was added to saturated brine, and extracted with t-butyl methyl ether. The organic layer was concentrated under reduced pressure,
20 and the residue obtained was subjected to silica gel column chromatography to give 155 mg of 5-cyclopentyl-3-(2-butynyloxy)-1,2,4-thiadiazole

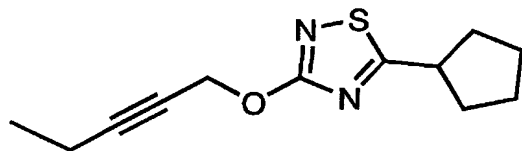
(hereinafter referred to as the present invention compound (3)).



$^1\text{H-NMR}$: 4.97 (s, 2H), 3.43 (m, 1H), 2.19 (m, 2H), 1.89-1.61 (m, 9H)

Production Example 4

5 By using 119 mg of 2-pentyn-1-ol instead of 2-butyn-1-ol according to Production Example 3 was obtained 187 mg of 5-cyclopentyl-3-(2-pentynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (4)).



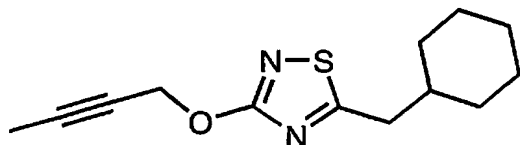
10 $^1\text{H-NMR}$: 5.00 (s, 2H), 3.45 (m, 1H), 2.23 (m, 4H), 1.87-1.68 (m, 6H), 1.13 (t, 3H)

Production Example 5

300 mg of 3-methylsulfonyl-5-(cyclohexyl)methyl-1,2,4-thiadiazole and 89 mg of 2-butyn-1-ol were dissolved in 2.5 g of N,N-dimethylformamide, 55 mg of sodium hydride (60% in oil) was added thereto, and the reaction mixture was stirred for 30 minutes under ice-cooling. Then, the reaction mixture was added to saturated brine, and extracted with t-butyl methyl ether. The organic layer was concentrated under reduced pressure, and the residue obtained was subjected to silica gel column chromatography to give 222 mg of 5-(cyclohexyl)methyl-3-(2-butynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (5)).

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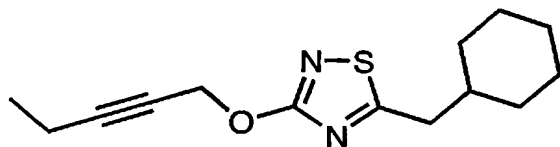
20



$^1\text{H-NMR}$: 4.98 (s, 2H), 2.89 (d, 2H), 1.87 (s, 3H), 1.82-1.62 (m, 6H), 1.31-0.98 (m, 5H)

Production Example 6

5 By using 106 mg of 2-pentyn-1-ol instead of 2-butyn-1-ol according to Production Example 5 was obtained 231 mg of 5-(cyclohexyl)methyl-3-(2-pentynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (6)).



10 $^1\text{H-NMR}$: 4.99 (s, 2H), 2.86 (d, 2H), 2.24 (m, 2H), 1.84-1.65 (m, 6H), 1.30-0.96 (m, 8H)

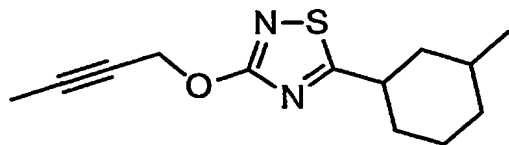
Production Example 7

350 mg of 3-methylsulfonyl-5-(3-methylcyclohexyl)-1,2,4-thiadiazole and 104 mg of 2-butyn-1-ol were dissolved in 3 ml of N,N-dimethylformamide, 75 mg of sodium hydride (60% in oil) was added thereto under ice-cooling, and the reaction mixture was stirred for 20 minutes under ice-cooling and for 20 minutes under room temperature. Then, the reaction mixture was added to saturated brine, and extracted with t-butyl methyl ether. The organic layer was concentrated under reduced pressure, and the residue obtained was subjected to silica gel column chromatography to give 280 mg of 5-(3-methylcyclohexyl)-3-(2-butynyloxy)-1,2,4-thiadiazole

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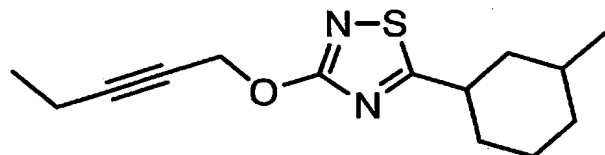
(hereinafter referred to as the present invention compound (7)).



¹H-NMR: 4.97 (s, 2H), 3.00 (m, 0.7H), 2.91 (m, 0.3H), 2.21-1.95 (m, 3H),
1.95-1.78 (m, 4H), 1.78-1.68 (d, 1H), 1.68-1.27 (m, 3H), 1.27-1.01 (m, 1H),
0.94 (m, 3H)

Production Example 8

By using 125 mg of 2-pentyn-1-ol instead of 2-butyn-1-ol according to
Production Example 7 was obtained 270 mg of 5-(3-methylcyclohexyl)-3-(2-
pentynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present
invention compound (8)).

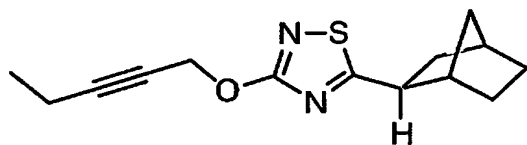


¹H-NMR: 4.99 (s, 2H), 3.00 (m, 0.7H), 2.90 (m, 0.3H), 2.23 (m, 2H), 2.14 (m,
2H), 2.06-1.68 (m, 3H), 1.68-1.28 (m, 3H), 1.13 (m, 4H), 0.95 (m, 3H)

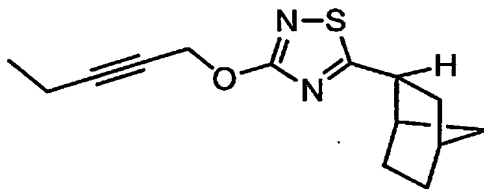
Production Example 9

300 mg of 3-methylsulfonyl-5-(2-norbornyl)-1,2,4-thiadiazole and 107
mg of 2-pentyn-1-ol were dissolved in 2.5 g of N,N-dimethylformamide, 60
mg of sodium hydride (60% in oil) was added thereto under ice-cooling, and
the reaction mixture was stirred for 30 minutes under ice-cooling and for 1
hour under room temperature. Then, the reaction mixture was added to
saturated brine, and extracted with t-butyl methyl ether. The organic layer
was concentrated under reduced pressure, and the residue obtained was

subjected to silica gel column chromatography to give 5-(2-norbornyl)-3-(2-butynyloxy)-1,2,4-thiadiazole. 150 mg of the *exo*-isomer having low polarity (hereinafter referred to as the present invention compound (9-1)) and 50 mg of the *endo*-isomer having high polarity (hereinafter referred to as the present invention compound (9-2)) were obtained by using hexane-ethyl acetate system as a carrier solvent.



¹H-NMR: 4.99 (s, 2H), 3.05 (t, 1H), 2.50 (s, 1H), 2.40 (s, 1H), 2.24 (m, 2H), 1.83 (d, 2H), 1.68-1.53 (m, 3H), 1.39 (m, 1H), 1.27 (m, 2H), 1.14 (t, 3H)

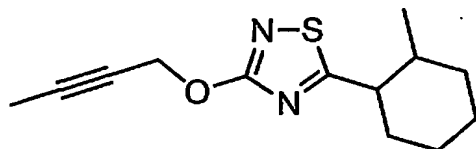


¹H-NMR: 4.99 (s, 2H), 3.45 (m, 1H), 2.66 (m, 1H), 2.38 (m, 1H), 2.23 (m, 2H), 2.11 (m, 1H), 1.55 (m, 2H), 1.45 (m, 2H), 1.39 (m, 1H), 1.26 (m, 2H), 1.16 (t, 3H)

Production Example 10

160 mg of 3-methylsulfonyl-5-(2-methylcyclohexyl)-1,2,4-thiadiazole and 52 mg of 2-butyne-1-ol were dissolved in 1.5 ml of N,N-dimethylformamide, 32 mg of sodium hydride (60% in oil) was added thereto under ice-cooling, and the reaction mixture was stirred for 1 hour under ice-cooling and for 12 hours at room temperature. Then, the reaction mixture was added to saturated brine, and extracted with t-butyl methyl ether. The organic layer was concentrated under reduced pressure, and the residue

obtained was subjected to silica gel column chromatography to give 39 mg of 5-(2-methylcyclohexyl)-3-(2-butynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (10)).

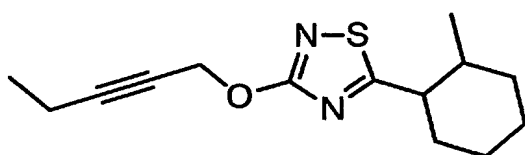


5 $^1\text{H-NMR}$: 4.98 (s, 2H), 3.05-2.93 (m, 1H), 2.16 (m, 2H), 1.88-1.65, 1.65-0.84 (m, 15H)

mass: $m/e=250(\text{M}^+)$

Production Example 11

By using 62 mg of 2-pentyn-1-ol instead of 2-butyne-1-ol according to
10 Production Example 10 was obtained 58 mg of 5-(2-methylcyclohexyl)-3-(2-pentyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (11)).

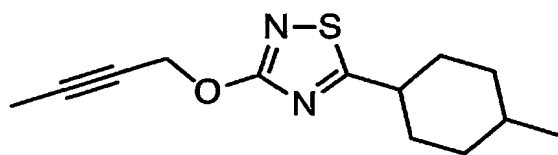


$^1\text{H-NMR}$: 5.05 (s, 2H), 3.04-2.93 (m, 1H), 2.30-0.84 (m, 17H)
15 mass: $m/e=264(\text{M}^+)$

Production Example 12

400 mg of 3-methylsulfonyl-5-(4-methylcyclohexyl)-1,2,4-thiadiazole and 129 mg of 2-butyne-1-ol were dissolved in 3 ml of N,N-dimethylformamide, 80 mg of sodium hydride (60% in oil) was added thereto
20 under ice-cooling, and the reaction mixture was stirred for 1 hour under ice-cooling and for 12 hours at room temperature. Then, the reaction mixture

was added to saturated brine, and extracted with t-butyl methyl ether. The organic layer was concentrated under reduced pressure, and the residue obtained was subjected to silica gel column chromatography to give 258 mg of 5-(4-methylcyclohexyl)-3-(2-butynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (12)).

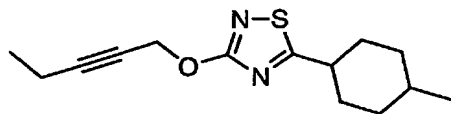


¹H-NMR: 4.98 (s, 2H), 3.02-2.85 (m, 1H), 2.15, 1.87-0.93 (m, 17H)

mass: m/e = 250 (M⁺)

Production Example 13

By using 155 mg of 2-pentyn-1-ol instead of 2-butyn-1-ol according to Production Example 12 was obtained 282 mg of 5-(4-methylcyclohexyl)-3-(2-pentynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (13)).



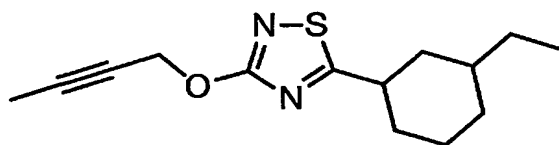
¹H-NMR: 5.00 (s, 2H), 3.05-2.86 (m, 1H), 2.23, 2.13, 1.83, 1.83-1.20, 1.20-1.00, 1.00-0.89 (m, 17H)

mass: m/e = 264 (M⁺)

Production Example 14

170 mg of 3-methylsulfonyl-5-(3-ethylcyclohexyl)-1,2,4-thiadiazole and 52 mg of 2-butyn-1-ol were dissolved in 1.5 ml of N,N-dimethylformamide, 32 mg of sodium hydride (60% in oil) was added thereto under ice-cooling, and the reaction mixture was stirred for 1 hour under ice-

cooling and for 12 hours under room temperature. Then, the reaction mixture was added to saturated brine, and extracted with t-butyl methyl ether. The organic layer was concentrated under reduced pressure, and the residue obtained was subjected to silica gel column chromatography to give
5 80 mg of 5-(3-ethylcyclohexyl)-3-(2-butynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present invention compound (14)).

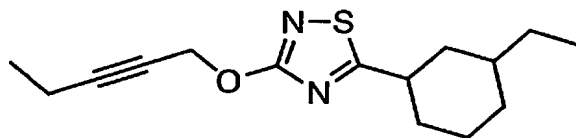


$^1\text{H-NMR}$: 4.97 (s, 2H), 3.03-2.90 (m, 1H), 2.16 (m, 2H), 1.95-1.78 (m, 6H),
1.48-1.25 (m, 5H), 1.13 (m, 1H), 0.90 (m, 3H)

10 mass: $m/e = 264 (M^+)$

Production Example 15

By using 63 mg of 2-pentyn-1-ol instead of 2-butyne-1-ol according to Production Example 14 was obtained 83 mg of 5-(3-ethylcyclohexyl)-3-(2-pentynyloxy)-1,2,4-thiadiazole (hereinafter referred to as the present
15 invention compound (15)).

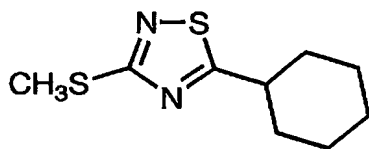


$^1\text{H-NMR}$: 4.99 (s, 2H), 3.03-2.93 (m, 1H), 2.28-2.13, 1.93-1.79, 1.46-1.23, 1.13,
0.89 (m, 19H)

20 Productions of intermediates of the present invention compounds are here exemplified as Reference Production Example.

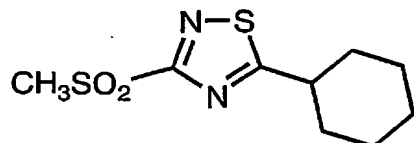
Reference Production Example 1

500 mg of 5-chloro-3-methylthio-1,2,4-thiadiazole and 122 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane complex were dissolved in 6 ml of tetrahydrofuran under nitrogen atmosphere, 6.58 ml of solution of cyclohexyl zinc bromide in tetrahydrofuran (0.5 mol/l) was added thereto under ice-cooling taking for about 5 minutes, and the reaction mixture was stirred for 16 hours at room temperature. Then, the reaction mixture was concentrated, and the residue subjected to silica gel column chromatography to give 370 mg of 3-methylthio-5-cyclohexyl-1,2,4-thiadiazole.



¹H-NMR: 3.05 (m, 1H), 3.68 (s, 3H), 2.13 (d, 2H), 1.86 (d, 2H), 1.74 (d, 1H), 1.50-1.24 (m, 5H)

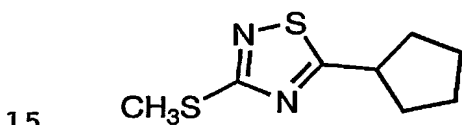
370 mg of 3-methylthio-5-cyclohexyl-1,2,4-thiadiazole was dissolved in 8 ml of chloroform, 1.12 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto, and the reaction mixture was stirred for 7 hours under ice-cooling. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 444 mg of 3-methylsulfonyl-5-cyclohexyl-1,2,4-thiadiazole.



¹H-NMR: 3.38 (s, 3H), 2.19 (m, 1H), 2.21 (d, 2H), 1.88 (d, 2H), 1.78 (d, 1H), 1.54-1.24 (m, 5H)

5 Reference Production Example 2

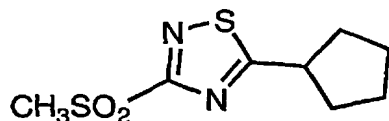
1.00 g of 5-chloro-3-methylthio-1,2,4-thiadiazole and 244 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane complex were dissolved in 12 ml of tetrahydrofuran under nitrogen atmosphere, 13.2 ml of solution of cyclopentyl zinc bromide in tetrahydrofuran (0.5 mol/l) was added thereto under ice-cooling taking for about 15 minutes, and the reaction mixture was stirred for 13 hours at room temperature. Then, the reaction mixture was concentrated, and the residue subjected to silica gel column chromatography to give 710 mg of 3-methylthio-5-cyclopentyl-1,2,4-thiadiazole.



¹H-NMR: 3.49 (m, 1H), 2.67 (s, 3H), 2.22-2.19 (m, 2H), 1.83-1.72 (m, 6H)

710 mg of 3-methylthio-5-cyclopentyl-1,2,4-thiadiazole was dissolved in 15 ml of chloroform, 1.89 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto, and the reaction mixture was stirred for 30 minutes under ice-cooling and for 3.5 hours at room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium

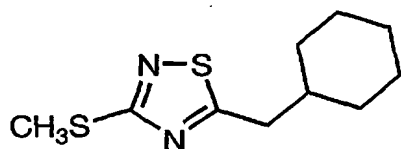
hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 804 mg of 3-methylsulfonyl-5-cyclopentyl-1,2,4-thiadiazole.



5 $^1\text{H-NMR}$: 3.61 (m, 1H), 3.37 (s, 3H), 2.31-2.26 (m, 2H), 1.88-1.77 (m, 6H)

Reference Production Example 3

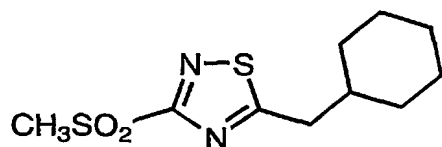
1.00 g of 5-chloro-3-methylthio-1,2,4-thiadiazole and 244 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane
10 complex were dissolved in 12 ml of tetrahydrofuran under nitrogen atmosphere, 13.2 ml of solution of (cyclohexyl)methyl zinc bromide in tetrahydrofuran (0.5 mol/l) was added thereto under ice-cooling taking for about 10 minutes, and the reaction mixture was stirred for 30 minutes under ice-cooling and for 8 hours at room temperature. Then, the reaction
15 mixture was concentrated, and the residue subjected to silica gel column chromatography to give 1.29 g of 3-methylthio-5-(cyclohexyl)methyl-1,2,4-thiadiazole.



20 $^1\text{H-NMR}$: 2.93 (d, 2H), 2.67 (s, 3H), 1.77-1.65 (m, 6H), 1.27-1.01 (m, 5H)

1.2 g of 3-methylthio-5-(cyclohexyl)methyl-1,2,4-thiadiazole was dissolved in 20 ml of chloroform, 2.78 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto, and the reaction mixture was stirred for

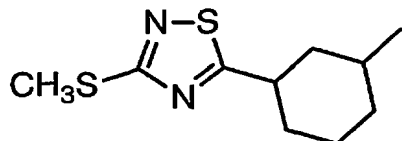
1 hour under ice-cooling and for 16 hours at room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 1.4 g of 3-methylsulfonyl-5-(cyclohexyl)methyl-1,2,4-thiadiazole.



$^1\text{H-NMR}$: 3.38 (s, 3H), 3.06 (d, 2H), 1.88-1.65 (m, 6H), 1.29-1.05 (m, 5H)

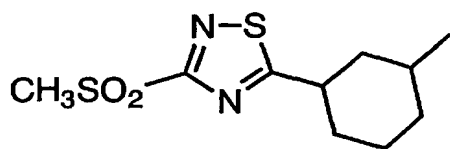
Reference Production Example 4

1.31 g of zinc foil (99.998%) and 190 mg of 1,2-dibromoethane was added into 2 ml of tetrahydrofuran under nitrogen atmosphere and the mixture was heated in an oil bath at 65°C for 1 minute. After cooling to room temperature, 0.1 ml of trimethylsilyl chloride, 2.14 g of 3-methylcyclohexyl iodide and 10 ml of tetrahydrofuran were added to the mixture, and the mixture was exposed to ultrasonic wave in a water bath for 4 hours. Further, 1.31 g of 5-chloro-3-methylthio-1,2,4-thiadiazole, 390 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane complex and 10 ml of tetrahydrofuran were added in the mixture, and the reaction mixture was stirred for 3.5 days at room temperature. The reaction mixture was concentrated, and the residue subjected to silica gel column chromatography to give 790 mg of 5-(3-methylcyclohexyl)-3-methylthio-1,2,4-thiadiazole.



$^1\text{H-NMR}$: 3.12-2.91 (m, 1H), 2.68 (s, 3H), 2.16 (m, 2H), 1.91-1.72 (m, 2H), 1.60-1.34 (m, 4H), 1.23-1.02 (m, 1H), 0.97 (t, 3H)

700 mg of 5-(3-methylcyclohexyl)-3-methylthio-1,2,4-thiadiazole was dissolved in 6 ml of chloroform, 1.73 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto, and the reaction mixture was stirred for 30 minutes under ice-cooling and for 1.5 hours under room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 784 mg of 3-methylsulfonyl-5-(3-methylcyclohexyl)-1,2,4-thiadiazole.

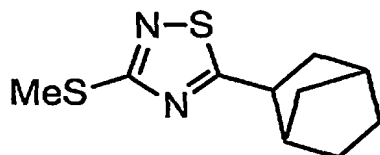


$^1\text{H-NMR}$: 3.38 (s, 3H), 3.24-3.02 (m, 1H), 2.24 (m, 2H), 1.96-1.74 (m, 2H), 1.67-1.38 (m, 4H), 1.28-1.06 (m, 1H), 0.98 (t, 3H)

Reference Production Example 5

1.67 g of 5-chloro-3-methylthio-1,2,4-thiadiazole and 245 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane complex were dissolved in 10 ml of tetrahydrofuran under nitrogen atmosphere, 21.2 ml of solution of *exo*-2-norbornyl zinc chloride in tetrahydrofuran (0.5 mol/l) was added thereto at room temperature, and the

reaction mixture was stirred for 1 day at room temperature. Then, the reaction mixture was concentrated, and the residue subjected to silica gel column chromatography to give 1.0 g of 5-(2-norbornyl)-3-methylthio-1,2,4-thiadiazole.



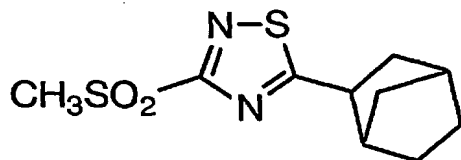
5

¹H-NMR: 3.50 (m, 0.3H), 3.15 (m, 0.7H), 2.68 (s), 2.67 (s, 3H), 2.51 (br), 2.42 (br), 2.18 (m), 1.95-1.79 (m), 1.70-1.33 (m), 1.28 (m, 10H)

1.0 g of 5-(2-norbornyl)-3-methylthio-1,2,4-thiadiazole was dissolved in 9 ml of chloroform, 3.67 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto, and the reaction mixture was stirred for 30 minutes under ice-cooling and for 1 day at room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 833 mg of 5-(2-norbornyl)-3-methylsulfonyl-1,2,4-thiadiazole.

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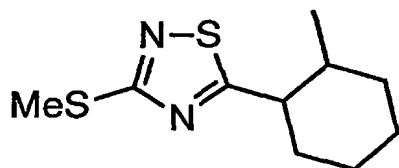
15



¹H-NMR: 3.63 (m, 0.3H), 3.38 (s, 0.3H), 3.37 (s, 0.7), 3.25 (m, 0.7H), 2.75 (m, 0.3H), 2.56 (br), 2.48 (br), 2.23 (m), 1.98 (m, 2H), 1.87 (m), 1.75-1.38 (m), 1.32 (m), 1.16 (m, 10H)

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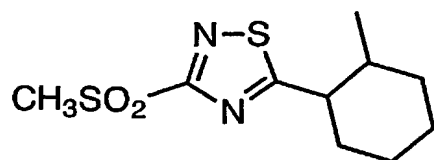
583 mg of zinc powder and 83 mg of 1,2-dibromoethane was added into 9 ml of tetrahydrofuran under nitrogen atmosphere and the mixture was refluxed for 1 minute. After cooling to room temperature, 49 mg of trimethylsilyl chloride and 3.0 g of 2-methylcyclohexyl iodide were added to the mixture, and the mixture was stirred for 6 hours at room temperature. Further, 1.49 g of 5-chloro-3-methylthio-1,2,4-thiadiazole and 109 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane complex were added in the mixture, and the reaction mixture was stirred for 15 hours at room temperature and refluxed for 5 hours. The reaction mixture was concentrated, and the residue subjected to silica gel column chromatography to give 564 mg of 5-(2-methylcyclohexyl)-3-methylthio-1,2,4-thiadiazole.



¹H-NMR: 3.12-3.02, 2.95 (m, 1H), 2.68 (s, 3H), 2.14 (m, 2H), 1.90-1.02 (m), 0.96 (m), 0.82 (m, 10H)

500 mg of 5-(2-methylcyclohexyl)-3-methylthio-1,2,4-thiadiazole was dissolved in 6.5 ml of chloroform, 1.62 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto taking for 20 minutes, and the reaction mixture was stirred for 30 minutes under ice-cooling and for 1.5 hours under room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 466 mg of 3-

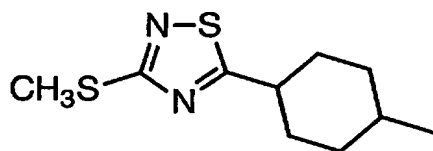
methylsulfonyl-5-(2-methylcyclohexyl)-1,2,4-thiadiazole.



¹H-NMR: 3.38 (m, 3H), 3.19 (m, 1H), 2.22 (br), 1.96-1.03 (m), 0.97 (m), 0.83 (m, 12H)

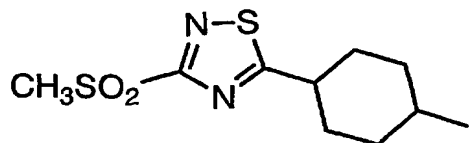
5 Reference Production Example 7

By using 3.0 g of 4-methylcyclohexyl iodide instead of 2-methylcyclohexyl according to Reference Production Example 6 was obtained 1.03 g of 5-(4-methylcyclohexyl)-3-methylthio-1,2,4-thiadiazole.



10 ¹H-NMR: 3.12-2.92 (m, 1H), 2.69 (s, 3H), 2.16 (m, 2H), 1.85 (m, 2H), 1.80-1.03 (m), 0.97 (m, 8H)

900 mg of 5-(4-methylcyclohexyl)-3-methylthio-1,2,4-thiadiazole was dissolved in 8 ml of chloroform, 2.22 g of 3-chloroperoxybenzoic acid (content >65%) was added thereto taking for 20 minutes, and the reaction mixture
15 was stirred for 4 hours at room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 1.11 g of 3-methylsulfonyl-5-(4-methylcyclohexyl)-1,2,4-thiadiazole.



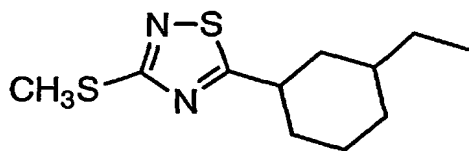
$^1\text{H-NMR}$: 3.39 (s, 3H), 3.27-3.05 (m, 1H), 2.26 (m), 2.14-1.06 (m), 0.97 (m, 12H)

Reference Production Example 8

5 439 mg of zinc powder and 63 mg of 1,2-dibromoethane was added into 7 ml of tetrahydrofurane under nitrogen atmosphere and the mixture was refluxed for 1 minute. After cooling to room temperature, 36 mg of trimethylsilyl chloride and 3.0 g of 3-ethylcyclohexyl iodide were added to the mixture, and the mixture was stirred for 6 hours at room temperature.

10 Further, 561 mg of 5-chloro-3-methylthio-1,2,4-thiadiazole and 137 mg of {1,1'-bis(diphenylphosphino)ferrocene} dichloro palladium(II) dichloromethane complex were added in the mixture, and the reaction mixture was refluxed for 17 hours. The reaction mixture was concentrated, and the residue subjected to silica gel column chromatography to give 384

15 mg of 5-(3-ethylcyclohexyl)-3-methylthio-1,2,4-thiadiazole.

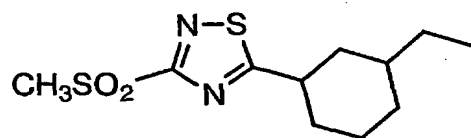


$^1\text{H-NMR}$: 3.06 (m, 1H), 2.68 (s, 3H), 2.22-2.14 (m, 2H), 1.92-1.80 (m, 2H), 1.45-1.09, 0.91 (m, 10H)

320 mg of 5-(3-ethylcyclohexyl)-3-methylthio-1,2,4-thiadiazole was dissolved in 2.5 ml of chloroform, 738 mg of 3-chloroperoxybenzoic acid (content >65%) was added thereto taking for 20 minutes, and the reaction

20

mixture was stirred for 20 minutes at room temperature. Then, the reaction mixture was added to saturated sodium hydrogensulfite aqueous solution, and separated. The organic layer was washed with sodium hydrogencarbonate aqueous solution, dried over anhydrous sodium sulfate, concentrated to give 380 mg of 3-methylsulfonyl-5-(3-ethylcyclohexyl)-1,2,4-thiadiazole.



$^1\text{H-NMR}$: 3.37 (s, 3H) , 3.23-3.09 (m, 1H) , 2.26 (m) , 1.95 (m) , 1.85 (m) , 1.58-1.02 (m, 11H) , 0.92 (t, 3H)

10

Formulation Examples are exemplified below. In addition, "part" means a part by weight. The present invention compounds are designated by their compound numbers shown above.

Formulation Example 1

9 Parts of each of the present invention compounds (1) to (15) are dissolved in 37.5 parts of xylene and 37.5 parts of dimethylformamide, and 10 parts of polyoxyethylene styryl phenyl ether and 6 parts of calcium dodecylbenzenesulfonate are added thereto, followed by well stirring and mixing, to give an emulsion for each compound.

Formulation Example 2

9 Parts of each of the present compounds (1) to (15) is added to a mixture containing 4 parts of sodium laurylsulfate, 2 parts of calcium lignin sulfonate, 20 parts of synthetic hydrated silicone oxide fine powder, and 65

parts of diatomaceous earth, followed by well stirring and mixing, to give a wettable powder for each compound.

Formulation Example 3

To 3 parts of each of the present invention compounds (1) to (15) are
5 added 5 parts of synthetic hydrated silicon oxide fine powder, 5 parts of sodium dodecylbenzenesulfonate, 30 parts of bentonite and 57 parts of clay, followed by well stirring and mixing. Then an appropriate amount of water is added to this mixture, followed by further stirring, granulating with a granulator and air drying, to give a granule for each compound.

10 Formulation Example 4

4.5 Parts of each of the present invention compounds (1) to (15), 1 part of synthetic hydrated silicon oxide fine powder, 1 part of Doriresu B (Sankyo Co., Ltd.) as a flocculant and 7 parts of clay are well mixed with a mortar, followed by stirring and mixing with a juice mixer. To the resulting
15 mixture is added 86.5 parts of cut clay, followed by well stirring and mixing, to give a powder for each compound.

Formulation Example 5

10 Parts of each of the present invention compounds (1) to (15), 35 parts of white carbon containing 50 parts of polyoxyethylene alkyl ether sulfate ammonium salt and 55 parts of water are mixed and pulverized by
20 the wet grinding method to give a formulation for each compound.

Formulation Example 6

0.5 Parts of each of the present invention compounds (1) to (15) are dissolved in 10 parts of dichloromethane, and the resulting solution is mixed
25 with 89.5 parts of Iso-Par M (isoparaffine: registered trade name for

EXXON CHEMICAL LTD) to give an oil solution.

Formulation Example 7

0.1 Parts of each of the present invention compounds (1) to (15) and 49.9 parts of NEO-CHIOZOL (CHUO KASEI Co., LTD) are charged into aerosol can, and aerosol valve is fixed to the can. Then 25 parts of dimethyl ether and 25 parts of LPG are filled in the can, followed by shaking and fitting an actuator on it, to give an oil aerosol.

Formulation Example 8

0.6 Parts of each of the present invention compounds (1) to (15), 0.01 parts of BHT, 5 parts of xylene, 3.39 parts of deodorized kerosene and 1 part of emulsifier [Atmos 300(registered trade name for ATMOS CHEMICAL LTD)] are mixed and dissolved. The resulting solution and 50 parts of distilled water are charged into aerosol container, and a valve is fixed to the container. 40 Parts of propellant (LPG) are charged under pressure through the valve to give an aqueous aerosol.

The following test example will demonstrate that the present invention compounds is useful as active ingredient of an arthropod pests controlling composition.

Test Example

The formulation of each of the present invention compounds (1) to (15) according to Formulation Example 5 was diluted with water so that the active ingredient concentration came to 500 ppm to prepare a spray solution.

A polyethylene cup was seeded with cucumber and a plant was grown until the first true leaf was developed, on which about twenty *Aphis gossypii*

(cotton aphid) are allowed to be parasitic. On the next day, the above spray solution was applied at a ratio of 20 ml/cup to the cucumber plant. On the sixth day after the application, the number of *Aphis gossypii* was examined.

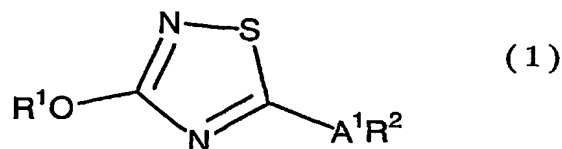
As a result, in the treatment with each of the present invention
5 compounds (1) to (15), the number of parasitic *Aphis gossypii* was not greater than 3.

Industrial Applicability

Arthropod pests such as insect pests, acarine pests and the like can
10 be effectively controlled by the present invention.

CLAIMS

1. A 1,2,4-thiadiazole compound represented by the formula (1):



- 5 wherein, R¹ represents C3-C7 alkynyl that may be substituted with halogen atom; R² represents C3-C8 cycloalkyl, C5-C8 cycloalkenyl or C7-C11 bicycloalkyl, wherein the C3-C8 cycloalkyl, the C5-C8 cycloalkenyl and the C7-C11 bicycloalkyl may be substituted with a substituent(s) selected from a group of C1-C4 alkyl, halogen atom and trifluoromethyl; A¹ represents a single
10 bond, C1-C2 alkylene or C2-C3 alkylidene.
2. The 1,2,4-thiadiazole compound according to claim 1, wherein R² is C3-C8 cycloalkyl which may be substituted with C1-C4 alkyl in the formula
15 (1).
3. The 1,2,4-thiadiazole compound according to claim 1 or 2, wherein A¹ is single bond in the formula (1).
4. The 1,2,4-thiadiazole compound according to claim 1 or 2, wherein A¹
20 is C1-C2 alkylene in the formula (1).
5. The 1,2,4-thiadiazole compound according to claim 1 or 2, wherein R¹ is 2-butynyl or 2-pentynyl in the formula (1).

6. An arthropod controlling composition comprising the 1,2,4-thiadiazole compound according to claim 1 and an inert carrier.
- 5 7. A method for controlling arthropod pests comprising applying an effective dose of the 1,2,4-thiadiazole compound according to claim 1 to the arthropod pests or habitat of the arthropod pests.
- 10 8. Use of the 1,2,4-thiadiazole compound according to claim 1 for the arthropod pests controlling composition.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/JP 03/13750

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D285/08 A01N43/836

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A01N A91N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, BEILSTEIN Data, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 30 30 661 A (BAYER AG) 1 April 1982 (1982-04-01) page 3 -page 4; claims 4,5	1-8
A	EP 0 273 534 A (NIPPON SODA CO) 6 July 1988 (1988-07-06) the whole document	1-8
A	EP 0 410 551 A (SCHERING AG) 30 January 1991 (1991-01-30) the whole document	1-8



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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Date of the actual completion of the international search

12 February 2004

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19/02/2004

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Fink, D

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/JP 03/13750

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